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Conti	nued Examination	RCE)	First	Named	Inventor		OBACI	H, R.S.	DF 20	1
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Request for Continu	t for Continued Examired Examination (RCE) practice gn application. See Instruct	tice under 37 CFR 1.11	l4 does no	at apply to	any util	ity or plan	nt application (	cation. iled prior t	o June 8.	
1. Submiss	ion required under 37	CFR 1.114								
а. 🗌	Previously submitted							•		
. •	Consider the amendment(s)/reply under 37 CFR 1.116 previously filed on									
ä	Consider the arg	uments in the Appeal 8	rief or Rel	y Brief pr	eviously	filed on _		-		
Ü.	Other: _	<del></del>	-							
b. 🔯	Enclosed									}
i.	Amendment/Rep	Ŋ	10.	$\boxtimes$	Inform	ation Dis	closure Stater	nent (IDS)	,	
ü.	Affidavit(s)/ Decl	aration(s)	iv.		Other			_		
2. Miscellar	neous					•				
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3. Fees	The RCE fee under 37 C	FR 1.17(e) is required	by 37 CFF	R 1.114 w	hen the	RCE is fil	ed.			
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i.	Deposit Account No RCE fee require	ed under 37 CFR 1.17(e	e)				•			
ii,	Extension of tim	e fee (37 CFR 1.136 and	1.17)							
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b. 🗀	Check in the amount of \$	enclosed								
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I hereby certify that this addressed to: Commission below.	a correspondence in baing depo sioner or Patents, Box RCE, Wi	CERTIFICATE OF MAI sited with the United State ishington, OC 20231, or fa	a Postal Se	vice with:	sufficient o	oostage as Patent and	first class mail i Trademark Offic	n an envelo	pe te	
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in the claims, the purpose of quinidine and ajmalacine is to inhibit CYP2D6, and not for "mediating oxidative biotransformation for the major clearance mechanism in humans" (Official Action, page 5, lines 4-6), as alleged by the Examiner. In contrast, (2S,3S)-2-phenyl-3-(2-methoxy-5-trifluoromethoxyphenyl)methylamino-piperidine is drug for which the major clearance mechanism in humans is CYP2D6 mediated oxidative biotransformation, and therefore has a different purpose from quinidine and ajmalacine. Thus, the instant case is distinguishable from In re Kerkhoeven, and therefore In re Kerkhoeven is not applicable to this case.

Applicants also submit a Declaration under 37 C.F.R. 132 by the inventor, Ronald Scott Obach, together with the data enclosed. As stated in the Declaration, in the enclosed data, Tables 1-4 describe enzymatic kinetic parameters for the metabolism of (2S,3S)-2-phenyl-3-(2-methoxy-5-trifluoromethoxyphenyl)methylamino-piperidine (including O-demethylation and N-dealkylation) in various mammals, and Table 5 describes the inhibition of the same compound by Cytochrome P450 isoform specific inhibitors. In the figures, Figures 10 and 11 show a correlation between metabolism and inhibition of the same compound using inhibitors quinidine (Figure 10) and ketoconazole (Figure 11). The foregoing data and figures show a surprising effectiveness of (2S,3S)-2-phenyl-3-(2-methoxy-5-trifluoromethoxyphenyl)methylamino-piperidine in combination with a CYP2D6 inhibitor such as, for example, quinidine or ketoconazole, further supporting the non-obvlousness of the invention over the cited art.

In view of the foregoing, withdrawal of the rejection of Claim 1 under 35 U.S.C. 103(a) as allegedly obvious over Benet et al. and Hess (WO 96/14845) is respectfully requested.

In view of the amendments and remarks made herein, applicants respectfully solicit the issuance of a notice of allowance. If a telephone interview is deemed to be helpful to expedite the prosecution of the subject application, the Examiner is invited to contact applicant's undersigned attorney at the telephone number provided.

The Commissioner is hereby authorized to charge any fees required under 37 C.F.R. §§1.16 and 1.17 or to credit any overpayment to Deposit Account No. 18-1445.

Date: December 2003

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